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1. A method for treatment of a subject having a disease or condition associated with apoptosis, which comprises administering an effective amount of a 15-keto-prostaglandin compound represented by the following formula (I):

$$R_1$$
 R_1
 R_1
 R_1
 R_1
 R_2
 R_3
 R_4
 R_4

wherein W₁, W₂ and W₃ are carbon or oxygen atoms;

L, M and N are hydrogen, hydroxy, halogen, lower alkyl, lower alkoxy, hydroxy(lower)alkyl or oxo, wherein at least one of L and M is a group other than hydrogen, and the five-membered ring may have one or more double bond(s);

A is -CH₂OH, -COOH or its functional derivative;

B is $-CH_2-CH_2^2$ -, -CH=CH- or $-C\equiv C$ -;

 R_1 is a divalent saturated or unsaturated lower-medium aliphatic hydrocarbon residue, which is unsubstituted or substituted by halogen, alkyl, hydroxy, oxo, anyl or heterocyclic group; and

Ra is a saturated or unsaturated lower-medium aliphatic hydrocarbon residue, which is unsubstituted or substituted by halogen, oxo, hydroxy, lower alkyl, lower alkoxy, lower alkanoyloxy, cyclo(lower)alkyl, cyclo(lower)alkyloxy, aryl, aryloxy, heterocyclic group or heterocyclic-oxy group; cyclo(lower)alkyl;

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cyclo(lower)alkyloxy; aryl; aryloxy; heterocyclic group; or heterocyclic-oxy group to the subject.

- The method of claim 1, wherein the 15-keto-prostaglandin 2. compound is a 13,14-dihydro-15-keto-prostaglandin compound.
- The method of claim 1, wherein the 15-keto-prostaglandin 3. 5 compound is a 15-keto-16-mono or dihalogen-prostaglandin compound.
 - The method of claim 1, wherein the 15-keto-prostaglandin 4. compound is a 13,14-dihydro-15-keto-16-mono or di-halogen-prostaglandin compound.
 - The method of claim 1, wherein the 15-keto-prostaglandin 5. compound is a 15-keto-16-mono or di-fluoro-prostaglandin compound.
 - The method of claim 1, wherein the 15-keto-prostaglandin 6. compound is a 13,14-dihydro-15-keto-16-mono or di-fluoro-prostaglandin compound.
 - The method of claim 1, wherein the 15-keto-prostaglandin 7. compound is a 15-keto-20-lower alkyl-prostaglandin compound.
 - The method of claim 1, wherein the 15-keto-prostaglandin 8. compound is a 15-keto-20-ethyl-prostaglandin compound.
- The method of claim 1, wherein the 15-keto-prostaglandin 9. compound is a 2-decarboxy-2-(2-carboxy lower alkyl)-15-keto-prostaglandin 20 compound.
 - The method of claim 1, wherein the 15-keto-prostaglandin 10. 2-decarboxy-2-(2-carboxyethyl)-15-keto-prostaglandin а compound is compound.
- The method of claim 1, wherein the 15-keto-prostaglandin 11. 25

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compound is a 2-decarboxy-2-(2-carboxyethyl)-13,14-dihydro-15-keto-16-mono or di-fluoro prostaglandin compound.

- The method of claim 1, wherein the 15-keto-prostaglandin 12. compound is a 2-decarboxy-2-(2-carboxyethyl)-13,14-dihydro-15-keto-16-mono or di-fluoro-20-ethyl-prostaglandin compound.
- The method of claim 1, wherein the 15-keto-prostaglandin 13. compound is a 2-decarboxy-2-(2-carboxyethyl)-13,14-dihydro-15-keto-16,16difluoro-20-ethyl-prostaglandin compound.
- The method of claim 1, wherein the 15-keto-prostaglandin 14. compound is a 15-keto-prostaglandin E compound.
- The method of claim 1, wherein the 15-keto-prostaglandin 15. 2-decarboxy-2-(2-carboxyethyl)-13,14-dihydro-15-keto-16,16compound is difluoro-20-ethyl-prostaglandin E₁ isopropyl ester.
- The method of claim 1, wherein the disease or condition 16. associated with apoptosis is an eye disorder caused by light.
- method of claim 1, which comprises administering The 17. ophthalmically a composition comprising a 15-keto-prostaglandin compound formulated in a dosage form suitable for ophthalmic administration.

The method of claim 17, wherein said composition is formulated

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as eye drops.

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